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This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

- 1. (Currently amended) A chimeric polypeptide, said chimeric polypeptide comprising:
 - a) a first domain comprising extracellular or an intracellular portions of a G protein coupled receptor, and
 - b) at least a second domain, attached to the first domain, wherein said second domain is <u>a</u> naturally or non-naturally occurring hydrophobic <u>moietymoieties</u>,

wherein said first domain does not comprise a native extracellular ligand portion of said GPCR.

- 2. (Original) The chimeric polypeptide of claim 1, wherein said second or more domains are attached at either one end, at both ends, or at an internal position of said first domain.
- 3. (Currently amended) The chimeric polypeptide of claim 1, wherein said hydrophobic moiety is selected from the group consisting of: a lipid, an acyl or an amino acid.
- 4. (Currently amended) The chimeric polypeptide of claim 3, wherein said hydrophobic moiety is selected from the group consisting of: stearoylecter (C18), palmitoylate (C16), myristoyl (C124), lauryl (C12), capryl (C10), and capryloyl (C68), phospholipids, steroids, sphingosines, ceramides, octyl glycine, 2-cyclohexylalanine, or benzolylphenylalanine, wherein said hydrophobic moiety is attached to said chimeric polypeptide with amide bonds, sulfhydryls, amines, alcohols, phenolic groups, or carbon-carbon bonds.
- 5. Cancelled.
- 6. (Original) The chimeric polypeptide of claim 1, wherein said intracellular portion is selected from the group consisting of: the first intracellular loop or a fragment thereof, the second intracellular loop or a fragment thereof, the third intracellular loop or a fragment thereof, and the fourth intracellular domain or a fragment thereof, of said G-protein coupled receptor.

7.-9. Cancelled.

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10. (Original) The chimeric polypeptide of claim 6, where said intracellular portion is at least 3 contiguous amino acid residues.

- 11. (Original) The chimeric polypeptide of claim 6, wherein said intracellular portion is at least 5 contiguous amino acid residues.
- 12. (Original) The chimeric polypeptide of claim 6, wherein said intracellular portion comprises the third intracellular loop of the GPCR.
- 13. (Original) The chimeric polypeptide of claim 12, wherein said intracellular portion comprises at least 7 contiguous amino acid residues of the third intracellular loop.
- 14. (Currently amended) The chimeric polypeptide of claim 1, wherein said <u>first domain</u> comprises a PAR and said second domain comprises a <u>lipid moietyGPCR transmembrane</u> domain or a fragment thereof.
- 15. Cancelled.
- 16. Cancelled.
- 17. Cancelled.
- 18. Cancelled.
- 19. (Currently amended) The chimeric polypeptide of claim 18, wherein the G-protein coupled receptor or fragment thereof, is selected from the group consisting of a luteinizing hormone receptor, a follicle stimulating hormone receptor, a thyroid stimulating hormone receptor, a calcitonin receptor, a glucagon receptor, a glucagon-like peptide 1 receptor (GLP-1), a metabotropic glutamate receptor, a parathyroid hormone receptor, a vasoactive intestinal peptide receptor, a secretin receptor, a growth hormone releasing factor (GRF) receptor, protease-activated receptors (PARs), cholecystokinin receptors, somatostatin receptors, melanocortin receptors, ADP receptors, adenosine receptors, thromboxane receptors, platelet activating factor receptor, adrenergic receptors, 5-HT receptors, CXCR4, CCR5, chemokine receptors, neuropeptide receptors, opioid receptors, erythropoietin receptor, von Willebrand receptor, parathyroid hormone (PTH) receptor, vasoactive intestinal peptide (VIP) receptor, and collagen receptors.
- 20. 28. Cancelled.
- 29. (Original) A pharmaceutical composition comprising the chimeric polypeptide of claim 1 and a pharmaceutically acceptable carrier.

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- 30. Cancelled.
- 31. (Original) A kit comprising in one or more containers, the pharmaceutical composition of claim 29.
- 32.- 34. Cancelled.
- --35. (New) The chimeric polypeptide of claim 1, wherein said G-protein coupled receptor is a mammalian G-protein coupled receptor.
- 36. (New) The chimeric polypeptide of claim 4, wherein said hydrophobic moiety is palmitoyl.
- 37. (New) The chimeric polypeptide of claim 19, wherein said G-protein coupled receptor is a protease-activated receptor (PAR).
- 38. (New) The chimeric polypeptide of claim 37, wherein the protease-activated receptor is selected from the group consisting of PAR1, PAR2, and PAR4.
- 39. (New) The chimeric polypeptide of claim 12, wherein said intracellular portion comprises a sequence selected from the group consisting of SEQ ID NO: 1-16, 19-23, and 29.
- 40. (New) The chimeric polypeptide of claim 12, wherein said intracellular portion comprises a sequence selected from the group consisting of SEQ ID NO: 1-10, and 23.
- 41. (New) The chimeric polypeptide of claim 1, wherein the said G-protein coupled receptor is selected from the group consisting of CCKA, CCKB, SSTR2, and SubP receptors.
- 42. (New) The chimeric polypeptide of claim 3, wherein said hydrophobic moiety is a steroid.
- 43. (New) A chimeric polypeptide, said chimeric polypeptide comprising:
- a) a first domain comprising an intracellular portion of a protease-activated receptor (PAR), and
- b) a second domain, attached to the first domain, wherein said second domain is palmitate.--